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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
10/044,070	01/11/2002	Jay Nelson	48892-1	7323		
22504 7	590 02/13/2004		EXAMINER			
DAVIS WRIG 2600 CENTUR	GHT TREMAINE, LI	ZARA, JANE J				
1501 FOURTH		ART UNIT	PAPER NUMBER			
SEATTLE, W	A 98101-1688	1635				

DATE MAILED: 02/13/2004

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.		Applicant(s)				
Office Action Summary		10/044,070		NELSON ET AL.				
		Examiner		Art Unit				
		Jane Zara		1635				
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply								
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.  - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.  - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.  - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.  - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).  Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).								
Status								
1) Responsive to communic	ation(s) filed on <u>11 Ja</u>	anuary 2002.						
2a) ☐ This action is FINAL.	This action is <b>FINAL</b> . 2b)⊠ This action is non-final.							
, , ,	☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.							
Disposition of Claims								
4) ☐ Claim(s) 1-10 is/are pending in the application.  4a) Of the above claim(s) is/are withdrawn from consideration.  5) ☐ Claim(s) is/are allowed.  6) ☐ Claim(s) is/are rejected.  7) ☐ Claim(s) is/are objected to.  8) ☐ Claim(s) 1-10 are subject to restriction and/or election requirement.								
Application Papers								
9) The specification is object	ed to by the Examine	r.						
10)☐ The drawing(s) filed on is/are: a)☐ accepted or b)☐ objected to by the Examiner.								
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).								
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).  11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.								
Priority under 35 U.S.C. § 119								
<ul> <li>12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).</li> <li>a) All b) Some * c) None of:</li> <li>1. Certified copies of the priority documents have been received.</li> <li>2. Certified copies of the priority documents have been received in Application No</li> <li>3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).</li> <li>* See the attached detailed Office action for a list of the certified copies not received.</li> </ul>								
Attachment(s)		_						
1) Notice of References Cited (PTO-892 2) Notice of Draftsperson's Patent Draw			Interview Summary ( Paper No(s)/Mail Da					
Information Disclosure Statement(s) (     Paper No(s)/Mail Date		5) 🔲		ce of Informal Patent Application (PTO-152)				

## DETAILED ACTION

Claims 1-10 are pending in the instant application.

## Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- Claims 1-5, drawn to an assay for determining therapeutic activity of US28 receptor antagonists, classifiable in class 435, subclass 3.
- II. Claims 6-9, drawn to a method for treating atherosclerosis or restenosis, classifiable in class 514, subclasses 1, 44.
- III. Claims 6-9, drawn to a method for treating chronic rejection syndrome or graft versus host disease, classifiable in class 514, subclasses 1, 44.
- IV. Claim 10, drawn to a method for enhancing cellular migration, classifiable in class 424, subclass 9.1.

The distinct and different inventions listed above are further restricted to distinct inventions each comprising a single receptor ligand and a single receptor antagonist.

Please elect a receptor ligand and a receptor antagonist in the election of Groups II and III. The methods involving different and distinct receptor ligands and antagonists comprise biologically and functionally different and distinct inventions, and thus one does not render the other obvious. The methods involving each distinct and different receptor ligand and antagonist comprise steps which are not required for or present in the methods of the other groups. Thus, the operation, function and effects of these

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different methods are distinct and different from each other, and capable of supporting separate patents.

Please elect <u>a single antisense sequence</u> for Groups II and III, for the reasons set forth below:

Pursuant to 35 U.S.C. 121 and 37 C.F.R. 1.141, the nucleotide sequences listed in claim 9 are subject to restriction. As per M.P.E.P. 2434, "the Commissioner has partially waived the requirements of 37 C.F.R. 1.141 and will permit a reasonable number of such nucleotide or amino acid sequences to be claimed in a single application." Under this policy, in most cases, up to 1 (one) independent and distinct nucleotide OR amino acid sequence will be examined in a single application without restriction. Those sequences which are patentably indistinct from the sequence selected by the applicant will also be examined.

Claim 9 specifically claims nucleotide sequences encoding antisense targeting US 28, and these individual SEQ ID Nos. are listed in claim 9. Each of these antisense sequences is considered to be structurally independent, because each of these sequences has a unique nucleotide sequence, and each target a specific region of US28. Furthermore, a search of all the sequences claimed presents an undue burden on the Patent and Trademark Office to search and examine all of the recited sequences. In view of the foregoing, applicants are required to elect up to 1 claimed nucleotide sequence from the claim.

The inventions are distinct, each from the other because of the following reasons:

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Inventions I and II and III and IV are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions comprise biologically and functionally different and distinct Groups and thus one does not render the other obvious. The methods of Groups I and II and III and IV comprise steps which are not required for or present in the methods of the other groups: determining therapeutic activity of US28 receptor antagonists comprising obtaining, isolating and determining smooth muscle cell migration in a migration device (Group I), treating atherosclerosis and restinosis in a subject (Group II), treating chronic rejection syndrome or graft versus host disease (Group III), stimulating a virally infected cell with a US28 receptor ligand (Group IV).

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Because these inventions are distinct for the reasons given above and have acquired a separate status in the art because of their recognized divergent subject matter, restriction for examination purposes as indicated is proper.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim

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remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

## Conclusion

Certain papers related to this application may be submitted to Art Unit 1635 by facsimile transmission. The faxing of such papers must conform with the notices published in the Official Gazette, 1156 OG 61 (November 16, 1993) and 1157 OG 94 (December 28, 1993) (see 37 C.F.R. § 1.6(d)). The official fax telephone number for the Group is 703-872-9306. NOTE: If Applicant does submit a paper by fax, the original signed copy should be retained by applicant or applicant's representative. NO DUPLICATE COPIES SHOULD BE SUBMITTED so as to avoid the processing of duplicate papers in the Office.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **Jane Zara** whose telephone number is **(571) 272-0765**. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, John LeGuyader, can be reached on (571) 272-0760. Any inquiry regarding this application should be directed to the patent analyst, Katrina Turner, whose telephone number is (571) 272-0564. Any inquiry of a general nature or relating to the status of this application should be directed to the Group receptionist whose telephone number is (703) 308-0196.

**JZ** February 5, 2004

SEAN MCGARRY PRIMARY EXAMINER

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